

Appl. No. 09, 89, 956
Amdt. dated August 6, 2003
Reply to Office action of March 10, 2003

REMARKS/ARGUMENTS

Claims 32-64 currently appear in this application. The Office Action of March 10, 2003, has been carefully studied. These claims define novel and unobvious subject matter under Sections 102 and 103 of 35 U.S.C., and therefore should be allowed. Applicants respectfully request favorable reconsideration, entry of the present amendment, and formal allowance of the claims.

Disclosure

The disclosure is objected to because there is no brief description of the drawings, it is not clear as to the composition of the activator and the coactivator, and there is a typographical error on page 14 for Bil.

Accordingly, the specification has been amended to correct these deficiencies.

With respect to the objections to the claims, claims 1-31 have been cancelled by the present amendment in favor of new claims 33-64.

Rejections under 35 U.S.C. 112

Claims 3, 5, 7, 10-15, 17, 19-21 and 24-29 are rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventors, at

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the time the application was filed, had possession of the claimed invention.

This rejection is respectfully traversed. The specification has been amended in accordance with the Examiner's helpful suggestions, incorporating the limitations of the claims into the specification. Since this information was part of the application as filed, it does not constitute new matter.

Claims 2, 3, 5, 7, 10, 14, 15, 17, 18, 20, 21, 26, 28 and 29 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject which applicant regards as the invention.

This rejection is respectfully traversed. Claims 1-31 have now been replaced by new claims 32-64. It is believed that claims 32-64 conform to the requirements of 35 U.S.C. 112.

Art Rejections

Claims 23, 26 and 27 are rejected under 35 U.S.C. 102(b) as being anticipated by Hase et al. Examples 9-14 of Hase et al. are said to teach cubic zinc sulfide electroluminophores having an average particle size of 8 or 9 microns.

This rejection is respectfully traversed. The compounds disclosed in Hase et al. are not

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electroluminophores, but are merely luminescent materials which exhibit a long lasting after-glow after being excited with electron beams. This is not the same as electroluminophores, which are used to manufacture electroluminescent lamps that are used either for background lighting of LCD displays or as luminous and marking elements. The luminescent materials described in Hase et al. are completely unsuitable for use as electroluminophores, and there is no indication that these compounds possess any electroluminescence.

Claims 23, 26 and 27 are rejected under 35 U.S.C. 102(b) as being clearly anticipated by Marking et al. or Mano.

This rejection is respectfully traversed. Contrary to the Examiner's assertion, Marking et al. disclose silver-activated zinc sulfide luminophores with low particle agglomeration. These luminescent materials are used in electron-beam tubes, and they are in no way able to provide efficient electroluminescence. In Marking et al., only silver is used as an activator for the zinc sulfide matrix. However, all effective zinc sulfide electroluminophores of the present invention are activated with the help of copper and/or gold.

The zinc sulfide electroluminophores of the present invention have a cubic crystal structure.

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However, the zinc sulfide crystals produced by Mano are baked to form a hexagonal crystal structure (column 6, lines 49-52), and then the crystals are milled to form crystal defects within the crystals, after which they are baked so that part of the whole crystal of the intermediate fluorescent material transited from a hexagonal system to a cubic system (column 7, lines 20-23, emphasis added). That is, Mano does not use crystals which are entirely cubic in conformation, but which are only partly cubic in conformation. The crystals made by the method of the present invention are wholly cubic. Therefore, it is respectfully submitted that Mano does not anticipate the zinc sulfide cubic crystals claimed herein.

The method of the present invention produces zinc sulfides with very narrow grain-size distributions without the requirement for sifting or harmful milling processes. Additionally, no steps need be taken to convert hexagonal crystals to cubic crystals, as the zinc sulfide crystals formed by the process of the present invention are all cubic.

Claim 22 is rejected under 35 U.S.C. 103(a) as being unpatentable over Mano. The Examiner states that Mano teaches cubic zinc sulfide electroluminophores having an average particle size of 20 microns or less.

This rejection is respectfully traversed. It is appreciated that Mano teaches zinc sulfide electroluminophores having an average particle size of 20 microns or less. However, Mano teaches that the hexagonal crystals must be subjected to milling so that part of the whole crystal is converted from hexagonal to cubic shape. This is not the same as zinc sulfide crystals which are cubic, and one skilled in the art reading Mano would not be motivated to produce zinc sulfide crystals which are completely cubic in structure, since Mano requires a milling step to convert part of the crystals to a cubic structure. In the present case, the method of making the crystals produces crystals which are different from those produced by Mano, particularly since the crystals of the present invention need not be milled to introduce strains into the crystals. Crystals with strains introduced therein cannot possibly be the same as crystals produced in the absence of introduced strains.

Claims 24 and 25 are rejected as being unpatentable over Hase et al. Hase et al. are said to teach that the phosphor can be subjected to any known surface treatment, and at page 13, it is said to teach that it is known in the art to apply protective layers.

This rejection is respectfully traversed. As noted above, Hase et al. do not disclose

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electroluminophores, but merely fluorescent particles.

The properties of these two types of particles are quite different. Therefore, one skilled in the art would have no motivation to apply the protective coatings of Hase et al. to the electroluminophore particles of the present invention.

Allowable Subject Matter

Claims 1, 8 and 16 would be allowable if rewritten or amended to overcome the objections set forth in this Office Action. Claims 2, 3, 5, 7, 10-15 and 17-21 would be allowable if rewritten or amended to overcome the rejections under 35 U.S.C. 112 set forth in this Office Action.

Accordingly, claims 1-31 have been rewritten as new claims 32-64. It is believed that new claims 32-64 conform to all of the requirements of 35 U.S.C. 112.

It is noted that the claimed process is not taught or suggested by the cited art of record. The Examiner states that there is no teaching of producing a zinc sulfide electroluminescent phosphor containing at least two activators, where one is not a halide, by the claimed method.

In view of the above, it is respectfully submitted that the claims are now in condition for

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allowance, and favorable action thereon is earnestly
solicited.

Respectfully submitted,

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Zinc Sulfide Electroluminophores and Methods for Their Production

REFERENCE TO RELATED APPLICATIONS

[0001] The present application is the national stage under 35 U.S.C. §371 of international application PCT/EP00/11069, filed November 9, 2000 which designated the United States, and which application was not published in the English language.

[0002] The present invention relates to zinc sulfide (ZnS) based fine-grain electroluminescent phosphors and methods for their production.

[0003] Phosphors of this type are generally doped with copper (Cu), however, optionally also with copper and/or gold (Au), as well as with copper and manganese (Mn) and they furthermore contain one or more coactivators, for which purpose halide anions (Cl, Br, I) or certain trivalent cations (e.g., Al, Ga, In) are usually inserted into the ZnS lattice.

[0004] In dependence upon their specific chemical composition, the zinc sulfide electroluminophores emit in the blue, green or yellow-orange range of the visible spectrum. In technical applications they are used to manufacture electroluminescent lamps that are used either for the background lighting of LCD displays (clocks, pocket calculators, mobile telephones, instrument lighting, etc.)

or as luminous and marking elements, e.g., in airplanes and motor vehicles, in the interior and on the facades of buildings, as well as in advertising fixtures, etc.

[0005] It is known that zinc sulfide electroluminophores, as compared to technical luminescent substances for the LTV, X-ray and cathode ray excitation, have a comparatively short life. The half-life (which is the time in which the brightness of the EL phosphors decreases to half its original value) of unencapsulated electroluminophores is only a few hundred hours. It is widely believed that their service life is significantly influenced, among other factors, by the grain size of the electroluminophores. This is one of the reasons why commercially available ZnS EL phosphors typically have average grain sizes in the range of 20 to 40 μm .

[0006] However, the extent to which pigments of such a large size can be made into high-quality layers is limited. In the commonly used screen printing techniques, for example, they require the use of comparatively large screens, resulting in dry layer thicknesses of up to 40 μm . These often display a visibly uneven EL emission due to the unavoidable inhomogeneities of such layers.

[0007] Another shortcoming of the resulting thick electroluminescent arrangements that is attributable to the large grain size of commercially available EL pigments lies

in the fact that relatively high supply voltages are required to attain the desired levels of brightness. These can result in high stresses on the incorporated binding agents and thus in a reduction in the service life of the EL arrangements.

[0008] Moreover, with the use of the coarse EL pigments according to the prior art, it is possible that individual particularly large luminophore particles can protrude from the layer despite the adjusted layer thickness of up to 40µm. In these cases the voltage stability and/or electric strength of the EL films is reduced, resulting in an additional reduction in their service life.

[0009] A significant reduction in the average grain size of the EL phosphors and simultaneous preservation or improvement of the brightness and durability values is therefore highly desirable for many technical applications that are based on the use of screen printing processes.

[0010] If, as recently proposed in DE 19 708 543, EL pigments are even to be processed into fine graphic structures, such as security elements in the field of value product printing by means of intaglio or offset printing processes, the availability of fine-grain EL phosphors must be considered an essential prerequisite for the technical feasibility of such an application. Experience has shown that it is necessary in this case to use average pigment

grain sizes of 2 to 6 gm to be able to meet the technical requirements of these printing processes.

[0011] Methods for producing efficient EL phosphors have been known for a long time. The pertinent prior art is described, for example, in U.S. Patent No. 4,859,361 and in WO 91/16722. According to those patent documents, the following steps are required to produce Cu doped or Cu and Mn doped ZnS electroluminophores that are coded with the usual coactivators:

Step 1: Preparation of a mixture of ZnS, the desired quantity of an EL activator (e.g., CuSO_4) and a coactivating, halide-containing fluxing agent (usually BaCl_2 , MgCl_2 , NaCl).

Step 2: Firing of this mixture at temperatures between 1000 and 1300°C .

Step 3: Cooling of the fired material to room temperature and rinsing with water.

Step 4: Mechanical working of the material by milling.

Step 5: Renewed firing of the thusly treated material in the temperature range between 600 and 900°C , optionally after previous renewed addition of ZnSO_4 and CuSO_4 .

Step 6: Cooling to room temperature and optional quenching with H_2O after a certain cooling time.

Step 7: Optional washing with H_2O and/or mineral acids to remove soluble components and with KCN solution to remove excess Cu_2S .

[0012] Particular importance is attached by the invention to the 4th preparation step. The mechanical working of the material that was first fired at 1000 to 1300°C is intended to transform a portion of the hexagonal ZnS electroluminophore formed under these conditions into the cubical crystal form. It is alleged that a transformation of this type effects an improvement of the brightness of the EL phosphors and particularly increases their life.

When the described process and comparable process variations are used, zinc sulfide electroluminophores with average grain sizes between 20 and 40 μm are obtained and individual particles may still significantly exceed this range of grain sizes. This can be attributed mostly to the high firing temperatures, as well as to the use of fluxing agents with a strongly mineralizing effect. Electroluminophores of this grain size class have the above-described shortcomings.

[0013] In patent document US 5,643,496, the process is modified to the extent that zinc sulfide electroluminophores can be obtained that have a grain size smaller than 23 μm , preferably 21 μm , and which, by adjusting the temperature of the first firing process to between 1100 and 1190°C, preferably to 1160°C, allegedly

reach the level of 25 μm large ZnS electroluminescent materials regarding their attainable levels of brightness and half lives.

[0014] Such a minor reduction in the average grain size of the electroluminophores hardly results in any noticeable improvements even for the use in screen printing processes. The principal shortcomings of EL phosphor particles of this coarseness largely still exist.

[0015] Average EL pigment grain sizes in the range of 10 μm can allegedly be attained with a process according to US 5,635,111; however, the solution described in that patent document has significant technical shortcomings. These lie in the fact that, on one hand, the firing is performed in a complicated vacuum apparatus in the presence of extremely aggressive and toxic gases (halogen halides, H_2S), which involves the associated risks if the apparatus should fail. On the other hand, the very time consuming and expensive process hardly appears suitable for producing larger quantities of EL luminophores under technical conditions. The present invention is therefore based on the object of creating a novel, cost-effective method for producing fine-grain zinc sulfide electroluminophores that can be made into efficient and long-lived electroluminescent layers of a high quality with various printing techniques.

SUMMARY OF THE INVENTION

[0016] In accordance with the present invention this object is met with the technical teaching, of claim 1. The inventive process is characterized accordingly by the following preparation steps and measures:

Step 1: Preparation of special, fine-grain zinc sulfides and use of these materials as a starting product for the synthesis of the inventive fine-grain zinc sulfide electroluminophores.

[0017] The preparation of zinc sulfide starting materials of this type takes place by precipitation of ZnS from the solutions of zinc salts, such as e.g., ZnSO_3 , $\text{Zn}(\text{NO}_3)_2$ and ZnCl_3 , preferably from zinc sulfate solutions, with the aid of induced H_2S gas or resulting from the addition of H_2S -generating compounds at temperatures of 20 to 80°C and a pH between 0.5 and 3.0. The zinc ion concentration of the given zinc salt solutions is adjusted to values of 0.25 moles/l to 2.0 moles/l.

[0018] This precipitation reaction produces as a result fine-grain zinc sulfides with very narrow grain-size distributions and the desired average grain size can be controlled by guiding the process parameters, such as, e.g., the zinc ion concentration, the speed with which the H_2S is passed in, the stirring speed, the temperature, and the pH.

[0019] The average grain sizes of the zinc sulfides prepared according to this invention and used as the

starting material for the synthesis of the inventive electroluminophores are typically 2 to 20 μm , preferably 2 to 5 μm or 5 to 15 μm .

[0020] This is a significant advantage compared to the prior art, because fine-grain zinc sulfides are already generated in this process step by purely preparative means, i.e., without the use of time-consuming sifting or harmful milling processes, and their grain size distribution is then transferred to the inventive electroluminophores by the specific design of the subsequent process steps and largely maintained throughout the sequential performance of these steps.

[0021] A further advantage lies in the fact that, in addition to the grain sizes, the surface properties of the ZnS precipitation products can also be controlled via the specific selected precipitation conditions. This results in compact ZnS crystallites with a very low tendency to form agglomerates. These special morphological characteristics of the ZnS starting materials prepared according to this invention advantageously affect the grain structure and the performance of the resulting inventive electroluminophores.

Step 2: Mixing of the fine-grain ZnS starting materials produced according to this invention with the activator-coactivator compounds required for the luminophore formation.

[0022] The copper and/or gold compounds, or copper and/or gold and manganese compounds (e.g. CuSO_4 , HAuCl_4 , $.4 \text{ H}_2\text{O}$, MnSO_4) which are used as activator materials, Au tetrachlorolaurate as well as aluminum compounds (e.g. $\text{Al}(\text{NO}_3)_3$) that may be required for the coactivation may already be added during the precipitation of the ZnS or also to the washed ZnS suspension after completion of the precipitation. This permits a homogeneous distribution of the activators and coactivators in the preparation mixture that is advantageous for the process of the luminophore formation, and also a close contact between the activator, coactivator and ZnS particles.

[0023] However, it is also possible to dry mix the activator and coactivator compounds and the zinc sulfide prepared according to this invention. In this case a preferred process variant consists of first homogenizing the activator and coactivator compounds with a portion of the dried ZnS and then blending this mixture with the remaining amount of ZnS that is required to ensure the desired luminophore composition.

[0024] To this mixture the fluxing agents, which are described in more detail below, are then added as well:

Step 3: A one to ten-hour firing of the mixture at temperatures below 1000°C , preferably in the temperature range between 800 and 1000°C in air or in an inert nitrogen

atmosphere or in an atmosphere consisting of a mixture of nitrogen and 1 to 10% hydrogen in the presence of fluxing agents with an only slightly mineralizing action selected from the compound classes of the fluorides, bromides and iodides.

[0025] After completion of the firing process, the fired product is then cooled to room temperature, subsequently washed with deionized water and then optionally filtered and dried.

[0026] In this manner it is effectively ensured that the average grain size and grain size distribution of the zinc sulfide electroluminophores after completion of the firing process and processing of the fired product essentially conform to the ZnS starting material prepared according to this invention and used for the luminophore synthesis. The inventive fine-grain ZnS electroluminophores that are obtained with this process step have typical medium grain sizes between 2 and 20 μm , preferably between 2 and 5 μm or 5 and 15 μm .

[0027] It is important compared to the prior art that the described firing process is performed at temperatures below 1000°C and that the presence, especially of chloride-containing or strongly mineralizing fluxing agents is completely abstained from. While it is true that the use of fluoride and/or bromide and/or iodide-containing fluxing

agents does enhance the reconstruction of the ZnS lattice and the targeted inclusion of the activators required for the formation of the luminophores, their operating mechanisms are such that the grain growth can effectively be limited in the described temperature range.

[0028] The fluxing agents that are used according to this invention can, at the same time, function as a source for the insertion of the coactivators. For this purpose they optionally receive certain trivalent cationic components (e.g. Al^{3+} Bi^{3+} in addition to the above halide anions and other cationic components).

[0029] A further advantage of the inventive process compared to the prior art lies in the fact that the synthesized electroluminophores remain completely in their cubic crystal modification because of the firing temperature being limited to a maximum of 1000°C . As will be described later, this fact results in advantages regarding the attainable levels of brightness and half-life of the inventive electroluminophores. In processes representing the prior art, hexagonally crystallizing ZnS electroluminophores are obtained initially. These are subsequently subjected to an intense and often harmful mechanical milling process in order to achieve an at least partial reverse transformation to the cubic structure type. The related shortcomings are prevented in advance with the present inventive process.

Step 4. Treatment of the powdery electroluminophores obtained after the main firing process with organic and inorganic acids.

[0030] The fine-grain zinc sulfide phosphors obtained after the implementation of preparation steps 1 through 4 are characterized by high photo and cathode luminescence yields. This fact is an indication of the effective inclusion of the activators and coactivators into the ZnS lattice as well as of the high effectiveness of the luminescence processes that occur under these excitation conditions.

[0031] It needs to be noted, however, that the phosphors that have been synthesized in this manner do not yet have optimum electroluminescent properties.

[0032] The efficiency of the electroluminescence can be increased significantly, however, if the zinc sulfide luminophore powder is subjected to a treatment with organic or inorganic acids, such as hydrochloric acid (HCl), sulfuric acid (H₂SO₄), nitric acid (HNO₃), acetic acid, or citric acid after the main firing process. For example, acid treatment can take place in about 37% HCl solution for two to six hours while stirring at room temperature. In another embodiment, acid treatment is with 10-20% citric acid for 4-8 hours at 60°C with stirring.

[0033] For this purpose the zinc sulfide phosphors obtained according to this invention are suspended in certain quantities of the solutions of these acids at temperatures between 20 and 60°C, preferably at room temperature, while stirring and the retention time of the luminophore particles in the given acid bath may span a range from 10 minutes up to 10 hours depending upon the type and concentration of the chosen acid, as well as on the chosen temperature. The powdery ZnS electroluminophores are subsequently filtered out, washed to pH neutrality and optionally dried at temperatures of 100 to 120°C.

[0034] As shown by electron-microscopic examinations, the inventive finest-grain electroluminophores show a clearly changed morphology of the crystallites after this acid treatment. It is characterized by a high roughness of the crystallite surfaces as well as by the formation of grooves, corners, edges and other structural defects. Such a modification of the habit of the doped ZnS crystallites apparently is an important prerequisite for the increase in the electroluminescence yields of the zinc sulfide luminophores that is noted after the acid treatment.

[0035] An advantageous secondary effect of the described acid treatment lies in the further reduction of the average grain size of the luminophore particles, as well as in the further narrowing of the grain size distributions. The scope

of this effect can be controlled via the conditions of the acid treatment. The acid treatment furthermore enhances the de-agglomeration of the luminophore particles, resulting in additional advantages for the use of the inventive luminophores in electroluminescent layers (dispersion behavior, layer homogeneity).

Step 5: Re-doping of the finest-grain electroluminophores that have been synthesized according to this invention with certain quantities of activator and/or coactivator ions.

[0036] Even with the inventive process it is possible to further increase the EL efficiency through the customary repeated addition of certain quantities of activator compounds, particularly Of CuSO_4 and/or coactivator compounds, particularly those containing Al^{3+} ions, and renewed firing of the resulting solids mixtures at temperatures between 300 and 800°C. In this manner a fine adjustment of the activator and/or coactivator concentrations and distribution of the active luminescence centers is achieved in the ZnS matrix.

[0037] This re-doping can be effected with copper, gold, manganese, and/or aluminum compounds. Preferred compounds for this re-doping include copper sulfate and/or tetrachloroauric acid or its sodium, salt, and/or manganese sulfate and/or aluminum nitrate.

[0038] Re-doping can take place in an air or an inert nitrogen atmosphere, or in an atmosphere consisting of a mixture of nitrogen and 1 to 10% hydrogen.

[0039] After a firing time of preferably 30 minutes to 10 hours, the fired product is cooled to room temperature and subsequently washed with H₂O, mineral acids (e.g., HNO₃), or KCN solution to remove activator and/or coactivator compounds that were not inserted into the ZnS lattice and have precipitated on the surface.

[0040] Alternatively, the ZnS is washed with a mineral acid and then with deionized water to pH neutrality and subsequently filtered and dried after their treatment with mineral acids of DCN solution following the re-doping.

Step 6: Annealing of the zinc sulfide electroluminophores obtained after preparation step 5 for 30 minutes to 5 hours at temperatures between 200 and 500°C.

[0041] Alternatively, this annealing step may take place after step 4. Annealing may take place in air or in an inert nitrogen atmosphere or in an atmosphere consisting of a mixture of nitrogen and 1 to 10% hydrogen.

[0042] This preparation step, which concludes the inventive process, serves for the final manifesting of the luminophore composition that is advantageous for the performance of the inventive finest-grain zinc sulfide electroluminophores.

[0043] The essence of the invention thus lies in combining the described process steps, particularly in carrying out the sequence of the first 4 steps listed.

BRIEF DESCRIPTION OF THE DRAWINGS

[0044] Figure 1 shows grain size distribution curves for zinc sulfide.

[0045] Figure 2 shows d_{50} values of grain size distribution of electroluminophores of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

[0046] With this method a fine-grain cubic zinc sulfide is already generated during the first step, which is then used as the starting material for the synthesis of the inventive electroluminophores, and the average grain size, grain size distribution and crystal structure of which is essentially maintained by limiting the firing temperature to a maximum of 1000°C and forgoing the use of fluxing agents with a strongly mineralizing effect in step 3 of the inventive process. At the same time the treatment of the luminophore powders obtained after the firing process with inorganic or organic acids according to step 4 together with the process steps 5 and 6 ensures that the electroluminophores synthesized according to this invention have all the composite and structural characteristics

required for attaining a high performance despite their small grain sizes.

[0047] The sequence of the above 4 process steps has now made it possible for the first time to obtain powerful fine-grain electroluminophores with grain sizes of 2 to 20 μm by purely preparative means and in a cost-effective manner without having to subsequently correct the grain size by means of milling, and screening, which would entail serious shortcomings.

[0048] For screen printing applications, inventive zinc sulfide electroluminophores with average grain sizes of 5 to 20 μm are used. Electroluminophores with these dimensions can be advantageously processed into high-performance EL lamps with a significantly improved layer structure.

[0049] Inventive fine-grain electroluminophores with average grain sizes between 2 and 5 μm , on the other hand, are particularly suitable for applications in intaglio or offset printing. They permit the implementation of fine electroluminescent graphic structures, e.g., as security marks in value product printing. These particles are particularly suitable for use in narrow-mesh screens of 120 meshes/inch.

[0050] In any case, the inventive luminophores are characterized by a brightness-service life relationship that is adapted to the given application and optimal with respect

to the adjusted grain size. Moreover, studies have confirmed that EL elements can be constructed using electroluminophores with average grain sizes of 6 μm produced according to this invention that display levels of brightness and half lives which, under identical operating conditions, are comparable to EL films manufactured from commercially available coarse-grained EL pigments with grain sizes of 20 to 40 μm .

[0051] As previously mentioned, the surprisingly high service life of the electroluminophores produced according to this invention, especially of those EL pigments that have average grain sizes of 2 to 5 μm in accordance with this invention, is very likely attributable to their cubic crystal structure, which deviates from the commercially available EL pigments.

[0052] According to the prior art this is considered advantageous for attaining high levels of brightness and stability.

[0053] The zinc sulfide electroluminophore particles can be coated with thin organic and/or crystalline or amorphous inorganic protective layers so as to increase their service life further. In one embodiment, the protective layer consists of an inner metal oxide film and an outer silicon nitrate film.

[0054] The zinc sulfide electroluminophore particles are dispersible, and can be used for printing in a halftone photogravure ink, flexographic printing ink, offset printing ink, letterset printing ink, gravure printing ink. The zinc sulfide electroluminophore particles of the present invention can be applied onto thermal transfer films and transferred to printable material by means of transfer printing. Alternatively, the zinc sulfide electroluminophore particles are embedded in thermoplastic granule matrices and processed into films by means of extrusion /coextrusion and/or thin film casting.

[0055] To further improve the stability, the individual crystallites of the inventive electroluminophores may also be provided with suitable protective layers according to the prior art. Numerous methods and materials are known for applying such protective layers.

[0056] Further details and advantages of the invention will be explained below based on examples and drawings.

Example 1

[0057] 10l of a 1.4 molar ZnSO_4 solution are entered into a reaction vessel. The pH of this solution is subsequently adjusted to 1.0 under addition of sulfuric acid (H_2SO_4).

[0058] The precipitation of the fine-grain zinc sulfide takes place by passing H_2S gas into the prepared solution while stirring (stirring speed 700 rpm). The volume flow of

the H₂S gas is 36 l/h, the work is performed at a reaction temperature of 60°C.

[0059] After a reaction time of approximately 500 min. the H₂S flow is stopped. Any H₂S still remaining in the reaction vessel is exhausted, the obtained ZnS suspension is decanted, repeatedly washed with deionized water and finally filtered off. The obtained fine-grain ZnS powder is subsequently dried at a temperature of 120°C.

[0060] Curve 1 in Figure 1 shows the grain size distribution of the fine-grain zinc sulfide prepared in this manner, which was determined with the aid of a Coulter counter grain size measuring instrument. What is striking is the very narrow distribution of the ZnS grain sizes (the so-called QD value, which is calculated based on the equation $QD = (d_{75} - d_{25}) / (d_{75} + d_{25})$, may be regarded as a measure for the range of the distribution, which, in the present case is $QD = 0.134$); a d_{50} value of 4.7 μm was determined for the average grain size of the ZnS material prepared according to the above described process.

[0061] In the next step a certain amount of the obtained zinc sulfide is stirred into a copper sulfate solution. After concentrating and drying of this suspension at approximately 120°C, the material, which is now present as a mixture of solids, is once again homogenized dry and subsequently sifted with a 35 μm gauze. The weighed-in

quantities of ZnS and CuSO₄ are calculated such as to establish a copper content of 1.5% for the zinc sulfide copper "activator".

[0062] A comparable process is also used in the preparation of corresponding ~~BiI₃~~-BiI₃ "activator". In the example described here, the ~~BiI₃~~-BiI₃ content of the ZnS-~~BiI₃~~-BiI₃ mixture is 8.5%.

[0063] The preparation of the batch subsequently takes place by a thorough blending of 1.65 kg of the fine-grain zinc sulfide, 81.5 g of the copper "activator", 7.5 g of the ZnS-~~BiI₃~~-BiI₃ mixture, as well as 5.2 g aluminum fluoride (AlF₃). This mixture is entered into covered quartz pans and fired for 2 hours at a temperature of 980°C in an N₂/H₂ atmosphere with a hydrogen content of 1.5%.

[0064] After completion of the firing process, the fired material is cooled to room temperature and repeatedly washed with deionized water.

[0065] This is followed by the acid treatment of the obtained material. For this purpose the washed fired material is entered into an acid bath and 2 l H₂O and 500 ml of a 37% hydrochloric acid are added relative to 1 kg of the fired material while stirring. After a retention time of one hour, this is followed by decanting and washing, with deionized water to pH neutrality.

[0066] The renewed addition of copper sulfate to this aqueous suspension serves to re-dope the luminophore material. The amount of CuSO_4 used for this purpose is calculated according to the ratio of 2 g Cu per 1 kg luminophore.

[0067] After concentration by evaporation and drying of the suspension, the dry material is fired in open quartz pans for 2 hours at 600°C in air. This is followed by an acid wash with 10% HNO_3 as well as repeated washing with H_2O to pH neutrality. This is followed by decanting, filtering and drying.

[0068] In a concluding process step, the obtained material is once again annealed in open quartz pans for 2 hours at 300°C in air and homogenized by sifting after it has cooled off.

[0069] As a result of these preparation steps a ZnS-Cu luminophore with a green electroluminescence is obtained that is characterized by a high level of brightness and long half-life. The average grain size of the powdery electroluminophore is $5.2\ \mu\text{m}$ ($\text{QD} = 0.265$). As can be seen from Fig. 1 (Curve 2), the average grain size of the EL pigment prepared according to the example is only significantly above that of the ZnS starting material used in this process.

Example 2

[0070] As in example 1, the precipitation of the zinc sulfide takes place after H_2S gas is passed into a ZnSO_4 solution, however, the reaction parameters are adjusted differently. The reaction is started with a 0.25 molar ZnSO_4 solution, the pH is fixed to 1.6, the H_2S volume flow is 60 l/h and the reaction temperature is 40°C .

[0071] The zinc sulfide that is present in the suspension after completion of the precipitation reaction has an average grain size of 17.0 μm ($\text{QD} = 0.174$, see Fig. 2, curve 1). The obtained ZnS suspension is washed repeatedly with deionized water and decanted; afterwards a sufficient amount of copper sulfate is added to establish a copper concentration of the ZnS material of 200 ppm after the activation. The copper activated ZnS suspension is transferred to drying pans and dried at 120°C .

[0072] To prepare the starting mixture for the firing process, 1.75 kg of the activated zinc sulfide, 0.5 g ~~BiI_3~~ **BiI_3** , and 2.5 AlF_3 are thoroughly blended. The firing takes place in covered quartz firing pans at 990°C in air. The firing time is 5 hours.

[0073] After cooling the fired product to room temperature and washing it with deionized water, a 5-hour acid treatment is performed with 20% citric acid. This is followed by decanting and washing with H_2O to pH neutrality.

[0074] The re-doping of the luminophore material again takes place through addition of copper sulfate (502.5 mg per 1 kg luminophore) to the aqueous ZnS:Cu suspension.

[0075] After concentration by evaporation and drying of the suspension the dry product is fired in open quartz pans for 3 hours at 700°C in air. This is followed by treatment with 10% HNO₃ and repeated washing with H₂O (to pH neutrality), decanting, filtering and drying.

[0076] The concluding annealing of the zinc sulfide electroluminophore takes place in open quartz pans for 1 hour at 500°C in air, followed by cooling and sifting.

[0077] The resulting ZnS:Cu luminophore has an intense blue electroluminescence, as well as a long half-life. As shown by Curve 2 in Fig. 2, the d₅₀ value of the grain size distribution of the inventive electroluminophore presented in this example, which describes the average grain size, is 14.5 μm (QD = 0.156) and thus somewhat below the value determined for the corresponding ZnS starting material.